

PATENT

Attorney Docket No. 067461-5041US06

(Former/Client Docket No.: A-67229-13)

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of

DAHIYAT *et al.*

Application No. 10/665,307

Filed: September 18, 2003

For: Protein Design Automation for
Protein Libraries

Examiner: DEJONG, Eric S.

Art Unit: 1631

Conf. No.: 6927

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Victoria Linne Poulsen

RESPONSE TO OFFICE ACTION

Mail Stop Amendment
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

This paper is being submitted in response to the Office Action mailed May 18, 2007. This response is timely filed on July 9, 2007. Although the Applicants do not believe any additional fees are required, the Commissioner is authorized to charge any additional fees, including extension fees or other relief, which may be required, or credit any overpayment to Deposit Account No. 50-0310 (Our order no. 067461-5041-US06 /67229-13/RMS).

Listing of the Claims begin on page 2.

Arguments/Remarks begin on page 7.

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Currently Pending Claims

This listing of claims will replace all prior versions and listings of claims in the application:

1. (Previously presented) A method for generating a secondary library of protein sequences of a target protein comprising:

- a) inputting the coordinates of said target protein into a computer;
- b) utilizing said coordinates and a forcefield calculation to generate a primary library of primary protein sequences comprising a plurality of primary variant amino acid residues at primary variant positions;
- c) generating a probability distribution of amino acid residues in a plurality of variant positions from said force field calculation;
- d) combining a plurality of said amino acid residues from said probability distribution to generate a secondary library of secondary sequences; wherein at least one of said secondary protein sequences is different from said primary protein sequences; and
- e) synthesizing a plurality of said secondary protein sequences.

2. (Previously presented) A method for generating a secondary library of protein sequences of a target protein comprising:

- a) inputting the coordinates of said target protein into a computer;
- b) utilizing said coordinates and a forcefield calculation to generate a primary library of primary protein sequences comprising a plurality of primary variant amino acid residues at primary variant positions;
- c) generating a probability distribution of amino acid residues in a plurality of variant positions from said force field calculation;
- d) generating a set of oligonucleotide probes each encoding at least one of said primary variant amino acid residues;
- e) using said probes in a polymerase chain reaction (PCR) to generate a plurality of oligonucleotide sequences, each encoding a plurality of secondary variant protein sequences, wherein at least one of said secondary variant protein sequences is different from said primary protein sequences; and,
- f) producing said secondary variant protein sequences in host cells transformed with said oligonucleotide sequences.

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3. (Previously Presented) A method according to claim 2 wherein said PCR is multiple PCR and wherein said probes are pooled.

4. (Previously Presented) A method according to claim 3 wherein said probes are added in equimolar amounts.

5. (Previously Presented) A method according to claim 3 wherein said probes are combined in amounts that correspond to the probability of said variant amino acid residues in said probability distribution table.

6. (Cancelled)

7. (Previously Presented) A method according to claim 1 wherein said target protein is an enzyme.

8. (Previously Presented) A method according to claim 1 wherein said target protein is a therapeutic protein.

9. (Previously Presented) A method according to claim 1 wherein the coordinates of a region surrounding a binding site to a receptor is input into a computer.

10. (Previously Presented) A method according to claim 1 wherein said primary variant positions comprise a region surrounding a binding site.

11. (Previously Presented) A method according to claim 8 wherein said primary variant positions comprise a region surrounding a binding site to a receptor.

12. (Previously Presented) A method according to claim 7 wherein said primary variant positions comprise a region surrounding the active site of said enzyme.

13. (Previously Presented) A method according to claim 7 wherein said primary variant positions comprise a region surrounding the catalytic residues of said enzyme.

14. (Currently Amended) A method for generating a secondary library of protein sequences of a target protein comprising:

(a) generating a primary library comprising:

(i) inputting the coordinates of a target protein with variable residue positions;

(ii) establishing a group of potential amino acids for each of said variable residue positions, wherein the group of potential amino acids for at least one of said variable residue position comprises at least two different amino acid side chains; and

(iii) analyzing the interaction of each of said amino acids with plurality of said amino acids at a plurality of variable residue positions and all or part of the remainder of said protein to generate a primary library of primary sequences;

(b) generating a probability distribution of amino acid residues from said primary library in a

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plurality of variant positions from said primary sequences;

(c) combining a plurality of said amino acid residues from said probability distribution to generate a secondary library of secondary sequences comprising secondary variants; wherein at least one of said secondary variants is different from said primary variants;[[,]] and

(d) synthesizing a plurality of said secondary protein sequences;

wherein at least one of said analyzing, generating or combining steps comprises using a force field calculation

15. (Currently Amended) A method for generating a secondary library of protein sequences of a target protein comprising:

(a) generating a primary library comprising:

(i) inputting the coordinates of a target protein with variable residue positions;

(ii) establishing a group of potential rotamers for each of said variable residue positions, wherein the group of potential rotamers for at least one of said variable residue position has a rotamer selected from each of at least two different amino acid side chains; and

(iii) analyzing the interaction of each of said rotamers with plurality of said rotamers at a plurality of variable residue positions and all or part of the remainder of said protein to generate a primary library of primary sequences;

(b) generating a probability distribution of amino acid residues from said primary library in a plurality of variant positions from said primary sequences;

(c) combining a plurality of said amino acid residues from said probability distribution to generate a secondary library of secondary sequences comprising secondary variants; wherein at least one of said secondary variants is different from said primary variants; and,

(d) synthesizing a plurality of said secondary protein sequences[[.]].

wherein at least one of said analyzing, generating or combining steps comprises using a force field calculation.

16. (Previously Presented) A method according to claim 14, wherein said force field calculation is Self-Consistent Mean Field (SCMF).

17. (Currently Amended) A method for generating a secondary library of protein variants of a target protein comprising:

(a) generating a primary library comprising:

(i) inputting the coordinates of a target protein with variable residue positions;

(ii) establishing a group of potential rotamers for each of said variable residue positions, wherein the group of potential rotamers for at least one of said variable residue position

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has a rotamer selected from each of at least two different amino acid side chains; and

- (iii) analyzing the interaction of each of said rotamers with plurality of said rotamers at a plurality of variable residue positions and all or part of the remainder of said protein to generate a primary library of primary sequences optimized for at least one scoring function;
- (b) generating a probability distribution of amino acid residues from said primary library in a plurality of variant positions from said primary sequences;
- (c) combining a plurality of said amino acid residues from said probability distribution to generate a secondary library of secondary sequences comprising secondary variants; wherein at least one of said secondary variants is different from said primary variants; and,
- (d) synthesizing a plurality of said secondary protein sequences[[]] .

wherein at least one of said analyzing, generating or combining steps comprises using a force field calculation.

18. (Previously Presented) A method according to claim 17; wherein said scoring function is selected from the group consisting of a van der Waals potential scoring function, a hydrogen bond potential scoring function, an atomic solvation scoring function, an electrostatic scoring function and a secondary structure propensity scoring function.

19. (Previously Presented) A method according to claim 14, 15, or 17, wherein said analyzing step utilizes a force field calculation.

20. (Previously Presented) A method according to claim 14, 15, or 17, wherein said generating step (B) utilizes a force field calculation.

21. (Previously Presented) A method according to claim 14, 15, or 17, wherein said combining step utilizes a force field calculation.

22. (Previously Presented) A method for generating a secondary library of protein sequences of a target protein comprising:

- a) inputting the coordinates of said target protein into a computer;
- b) specifying a list of at least two primary variant positions;
- c) utilizing said coordinates and a forcefield calculation to generate a primary library comprising a plurality of primary variant amino acid residues at said primary variant positions;
- d) generating a probability distribution of amino acid residues in a plurality of variant positions from a force field calculation;
- e) combining a plurality of said amino acid residues from said probability distribution to generate a secondary library of secondary sequences; wherein at least one of said secondary protein sequences is different from said primary protein sequences; and,

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f) synthesizing a plurality of said secondary protein sequences.

23. (Previously Presented) A method according to claim 22 wherein said primary library is generated using a monte carlo search.

24. (Previously Presented) A method according to claim 22 wherein said primary library is generated using a genetic algorithm.

25. (Previously Presented) A method according to claim 22 wherein said probability distribution table is derived from frequencies of occurrence in a primary variant library.

26. (Previously Presented) A method according to claim 1 wherein combining said plurality of said amino acid residues from said probability distribution comprises a calculation in said computer.

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REMARKS

Claims 1-5 and 7-26 are pending. Claims 14, 15 and 17 have been amended to correct a typographical error(s).

Claim Objections

Applicants appreciate the Examiner's indication that the objection to claims 2-5 is withdrawn.

Claim Rejection under 35 USC 101

Applicants appreciate the Examiner's indication that the rejection of claims 1 and 7-26 under 35 USC 101 is withdrawn.

Double Patenting

Claims 1-5 and 7-26 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 4-8 of US Patent No. 6403312. Submitted herewith is a terminal disclaimer obviating this rejection.

Claims 1-5 and 7-26 are provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 36-41 of copending Application No. 09/782,004. Applicants note the Examiner identified the instant application as a divisional of copending application 09/782,004. Nonetheless, the claims are provisionally rejected. Without necessarily agreeing with the propriety of the rejection and in no way acquiescing to the rejection, Applicants are submitting herein a terminal disclaimer obviating this rejection.

Claims 1-5 and 7-26 are provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 52-69 of copending Application No. 09/927,790. Submitted herewith is a terminal disclaimer obviating this rejection.

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Conclusion

Applicants believe the present application is in condition for allowance. Early favorable communication thereof is respectfully requested. Please direct any calls in connection with this application to the undersigned at (415) 442-1216.

While Applicants believe that no other fees are due at this time, the Commissioner is authorized to charge any fees, including extension fees or any other relief that may be required, in connection with this reply to Deposit Account 50-0310 (Docket No.: 67461-5041US06).

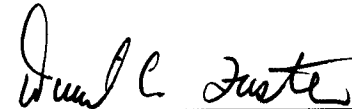
The undersigned is an attorney or agent of record.

MORGAN, LEWIS & BOCKIUS LLP

Dated: July 9, 2007

Customer No.: 67374
Morgan, Lewis & Bockius LLP
One Market, Spear Street Tower
San Francisco, CA 94105
Telephone: (415) 442-1000
Facsimile: (415) 442-1001

By:



David C. Foster, Reg. No. 44,685 for
Robin M. Silva, Reg. No. 38,304
Attorney of Record for Applicant
Filed Under 37 C.F.R. 1.34

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